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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO
10/740,698	12/19/2003	Signe Erickson Varner	SRM0061/US/2	3885
72870 7590 07/01/2011 Kagan Binder, PLLC			EXAMINER	
221 Main Stre		MEHTA, BHISMA		
Suite 200 Stillwater, MI	N 55082		ART UNIT	PAPER NUMBER
Summater, mi	. 55002		3767	
			MAIL DATE	DELIVERY MODE
			07/01/2011	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

## **Advisory Action** Before the Filing of an Appeal Brief

Application No.		Applicant(s)	
	10/740,698	VARNER ET AL.	
	Examiner	Art Unit	
	BHISMA MEHTA	3767	

	BHISMA MEHTA	3/6/					
The MAILING DATE of this communication appe	ars on the cover sheet with the c	orrespondence add	ress				
THE REPLY FILED 17 June 2011 FAILS TO PLACE THIS APPLICATION IN CONDITION FOR ALLOWANCE.							
<ol> <li>X The reply was filed after a final rejection, but prior to or on application, applicant must timely file one of the following application in condition for allowance; (2) a Notice of Appe for Continued Examination (RCE) in compliance with 37 C periods:</li> </ol>	replies: (1) an amendment, affidavit eal (with appeal fee) in compliance	, or other evidence, whith 37 CFR 41.31; or	hich places the (3) a Request				
a) The period for reply expires 3 months from the mailing date	of the final rejection.						
<ul> <li>The period for reply expires on: (1) the mailing date of this As no event, however, will the statutory period for reply expire la</li> </ul>	dvisory Action, or (2) the date set forth i ater than SIX MONTHS from the mailing	date of the final rejection	n.				
Examiner Note: If box 1 is checked, check either box (a) or (I MONTHS OF THE FINAL REJECTION, See MPEP 706.07(f		FIRST REPLY WAS FI	LED WITHIN TW				
Extensions of time may be obtained under 97 CFR 1.138(a). The date in have been filled is the date for purposes of determining the period of valued or 37 CFR 1.17(a) is calculated from: (1) the expiration date of the set forth in (b) above, if checked. Any reply received by the Office later may reduce any earned patient term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL.	ension and the corresponding amount of hortened statutory period for reply origin than three months after the mailing date	of the fee. The appropria nally set in the final Office	ate extension fee e action; or (2) as				
The Notice of Appeal was filed on A brief in compl filing the Notice of Appeal (37 CFR 41.37(a)), or any exter Notice of Appeal has been filed, any reply must be filed with the property of the p	sion thereof (37 CFR 41.37(e)), to	avoid dismissal of the					
<u>AMENDMENTS</u>							
<ol> <li>The proposed amendment(s) filed after a final rejection, be (a) They raise new issues that would require further core (b) They raise the issue of new matter (see NOTE below</li> </ol>	sideration and/or search (see NOT		cause				
(c) They are not deemed to place the application in bett appeal; and/or		lucing or simplifying ti	ne issues for				
(d) ☐ They present additional claims without canceling a c NOTE: (See 37 CFR 1.116 and 41.33(a)).	corresponding number of finally reje	cted claims.					
4. The amendments are not in compliance with 37 CFR 1.12	1. See attached Notice of Non-Cor	mpliant Amendment (	PTOL-324)				
5. Applicant's reply has overcome the following rejection(s):		. p. a					
<ol> <li>Applicant's reply has overcome are following rejection(s):        </li> <li>Co.  Newly proposed or amended claim(s):         would be allowable if submitted in a separate, timely filed amendment canceling non-allowable claim(s).</li> </ol>							
7.  For purposes of appeal, the proposed amendment(s): a) [how the new or amended claims would be rejected is prov The status of the claim(s) is (or will be) as follows:		be entered and an ex	xplanation of				
Claim(s) allowed:							
Claim(s) objected to:	•						
Claim(s) rejected: 68-74,76-119,122-127,129 and 132-140 Claim(s) withdrawn from consideration:	2-						
AFFIDAVIT OR OTHER EVIDENCE							
The affidavit or other evidence filed after a final action, but because applicant failed to provide a showing of good and was not earlier presented. See 37 CFR 1.116(e).							
<ol> <li>The affidavit or other evidence filed after the date of filing entered because the affidavit or other evidence failed to or showing a good and sufficient reasons why it is necessary</li> </ol>	vercome <u>all</u> rejections under appea	I and/or appellant fail:	s to provide a				
10. The affidavit or other evidence is entered. An explanation REQUEST FOR RECONSIDERATION/OTHER							
The request for reconsideration has been considered but See Continuation Sheet.	t does NOT place the application in	condition for allowan	ce because:				
12. Note the attached Information <i>Disclosure Statement(s)</i> . (13. Other:	PTO/SB/08) Paper No(s)						
	/Bhisma Mehta/ Primary Examiner, Art U	nit 3767					

Continuation of 11.: Applicant's arguments have been considered but are not deemed persuasive.

As to the arguments in line 13 of page 2 to line 23 of page 3, in Applicant's specification, the device has been disclosed as having deviations from a linear path. However, there is no discloser or support that the distinction between linear or non-linear is through following the "longitudinal axis" of the device. The claims are drawn to a non-linear shaped member and the outer surface of the body member of Weiner does provide a non-linear shape to the body member. The broadest reasonable interpretation consistent with the specification has been applied to the 'non-linear shaped body member' to indicate a body member which has a surface that does not follow a straight line such as one that has turn or angles. In lines 21-25 of page 7 of the specification, Applicant discloses that the device can have multiple turns or angles and the body member of Weiner does have multiple turns or angles. Even though Applicant states that "the overall shape of the capsule is linear" with regards to the device of Weiner, the body member of the different embodiments of the device of Weiner are not entirely linear and, thus, the shape of the devices are non-linear. As to the arguments in line 24 of page 3 to line 30 of page 4, Weiner only discloses that injection is preferred as a way of inserting the device into the obody. Weiner also discloses the device being inserted without injection where "a slight inviting orn" is specifiered during the insertion to facilitate entry. Modifying the device of Weiner to have a coil or zig-zag shape along its entire length would still enable one to insert the device with a slight withing most one as disclosed by Weiner.

As to the arguments in lines 1-25 of page 5, reducing the cross-sectional diameter would not considerably limit the amount of drug that could be placed in the device of Weiner as modifying the device of Weiner would still result in the device being of the same overall size such that approximately the same amount of drug could be placed in the device. Furthermore, one skilled in the art would recognize that the concentration of the drug in the device could be adjusted to correlate with the size of the device.

As to the arguments in line 26 of page 5 to line 8 of page 6, the primary reference of Welner teaches to device that is insertable through an incision and the device of Rosenman is indicated as being insertable or capable of being inserted through an incision. Trus, modification of the device of Welner to have a coil or 2jc-zag shape would still lead to a device that is insertable through an incision as claimed. As to the arguments in line 9 of page 6 to line 17 of page 8, Rosenman is not a dissimilar area of medical treatner as both Weiner and Rosenman disclose implantable devices and methods for inserting the implantable devices such that a drug or medicament can be delivered to a specific location in the patient's body.

As to Applicant's arguments in line 18 of page 8 to line 15 of page 9, the problem solved by the Rosenman technology does correlate to the problem solved of the current application as Rosenman discloses implantable devices of different shapes to allow for the delivery of a drug or medicament to a particular location in the patient's body. Furthermore, Weiner also discloses the use of implantable device of different shapes to deliver the drug to the desired location in the patient's body